CLAIMS

WE CLAIM:

- 1. A method of producing a human neural cell comprising,
 - a) providing a pluripotent human cell; and
 - b) culturing the pluripotent human cell with a composition comprising a ceramide compound of the general formula

wherein

R=a saturated or mono- or polyunsaturated (cis or trans) alkyl group having greater than 2 carbon atoms, and

R1, R2, R3 and R4 may be the same or different and are saturated or mono-or polyunsaturated hydroxylated alkyl groups, aryl groups, or hydrogen

to produce the human neural cell.

- 2. The method of Claim 1, wherein the pluripotent human cell is a differentiating pluripotent human cell.
- 3. The method of Claim 1, comprising the intermediate step of forming an embryoid body comprising the pluripotent human cell prior to culturing a cell from the embryoid body with the ceramide compound.
- 4. The method of Claim 3, wherein the embryoid body is formed by culturing the pluripotent human cell with an essentially serum free medium.
- 5. The method of Claim 4, wherein the essentially serum free medium is a MEDII conditioned medium.
- 6. The method of Claim 5, comprising the additional steps of,
 - a) dispersing the embryoid body to an essentially single cell suspension;

- b) culturing the essentially single cell suspension comprising the pluripotent human cell in an adherent culture with a composition comprising the ceramide compound.
- 7. The method of Claim 6, wherein the composition comprising the ceramide compound further comprises a MEDII conditioned medium.
- 8. The method of any of Claims 5, 6, or 7, wherein the MEDII conditioned medium is a Hep G2 conditioned medium.
- 9. The method of Claim 7, wherein the composition comprising the ceramide compound is essentially serum free.
- 10. The method of Claim 1, wherein the composition comprises a ceramide compound of the structure

11. The method of Claim 1, wherein the composition comprises a ceramide compound of the structure

- 12. The method of Claim 1, wherein the concentration of the ceramide compound is from approximately 0.1 μ M to approximately 1000 μ M.
- 13. The method of Claim 1, wherein the concentration of the ceramide compound is from approximately 1 μ M to approximately 100 μ M.
- 14. The method of Claim 1, wherein the concentration of the ceramide compound is from approximately 5 μ M to approximately 50 μ M.

15. The method of Claim 1, wherein the concentration of the ceramide compound is approximately 10 μ M.

- 16. The method of Claim 1, wherein the duration of culturing the human pluripotent cell with the ceramide compound is from approximately 6 hours to 10 days.
- 17. The method of Claim 1, wherein R= a saturated or mono- or polyunsaturated (cis or trans) alkyl group having from 12-20 carbon atoms.
- 18. The method of Claim 17, wherein the hydroxylated alkyl groups have from 1-6 carbon atoms.
- 19. The method of Claim 18, wherein R1 and R2 are hydroxylated alkyl groups.
- 20. The method of Claim 1, wherein the pluripotent human cell is selected from the group consisting of a human embryonic stem cell, a human inner cell mass (ICM)/epiblast cell, a human primitive ectoderm cell, and a human primordial germ cell.
- 21. The method of Claim 1, wherein the pluripotent human cell is a human embryonic stem cell.
- 22. The method of Claim 1, wherein the human pluripotent cell is a multipotent cell.
- 23. The method of Claim 22, wherein the multipotent cell is a neural precursor cell.
- 24. A method of producing a human neural cell comprising,
 - a) providing a pluripotent human cell;
 - b) forming an embryoid body comprising the pluripotent human cell by culturing the pluripotent human cell in a medium that is optionally essentially serum free; and
 - c) culturing cells from the embryoid body with a composition comprising a sphingosine compound of the general formula

wherein

R=a saturated or mono- or polyunsaturated (cis or trans) alkyl group

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having greater than 2 carbon atoms, and R1, R2, R3 and R4 may be the same or different and are saturated or mono-or polyunsaturated hydroxylated alkyl groups, aryl groups, or hydrogen.

- 25. The method of Claim 24, wherein the medium is an essentially serum free medium and comprises a MEDII conditioned medium.
- 26. A method of producing a human neural cell comprising,
 - a) providing a pluripotent human cell;
 - b) forming an embryoid body comprising the pluripotent human cell by culturing the pluripotent human cell in a medium that is optionally essentially serum free; and
 - c) culturing cells from the embryoid body with a composition comprising a hydroxyalkyl ester compound of the general formula

wherein

R=a saturated or mono- or polyunsaturated (cis or trans) alkyl group having greater than 2 carbon atoms, and R1 is a saturated or mono-or polyunsaturated hydroxylated alkyl group, aryl group, or hydrogen.

- 27. The method of Claim 26, wherein the medium is an essentially serum free medium and comprises a MEDII conditioned medium.
- 28. The method of Claim 27, wherein the composition comprises an hydroxyalkyl ester compound of the structure

29. A neural cell produced by any of the methods of Claims 1-28.

- 30. A method for treating a patient, comprising a step of administering to the patient having a neural disease a therapeutically effective amount of the neural cell of Claim 29.
- 31. The method of Claim 30, wherein the neural disease is Parkinson's disease.
- 32. A method of enhancing the efficiency of the transplantation of a cultured human pluripotent cell in a patient, comprising the steps of:
 - a) culturing a human pluripotent cell with a growth medium comprising a ceramide compound of the formula

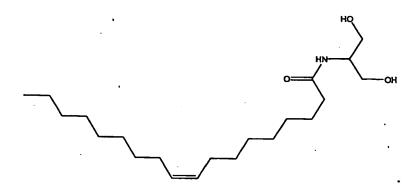
wherein

R=a saturated or mono- or polyunsaturated (cis or trans) alkyl group having greater than 2 carbon atoms, and

R1, R2, R3 and R4 may be the same or different and are saturated or mono-or polyunsaturated hydroxylated alkyl groups, aryl groups, or hydrogen; and

- b) transplanting the cultured human pluripotent cell into the patient.
- 33. The method of Claim 32, wherein the composition comprises a ceramide compound of the structure

34. The method of Claim 32, wherein the composition comprises a ceramide compound of the structure



- 35. The method of Claim 32, wherein the duration of step (a) is for a period of from approximately 6 hours to 10 days.
- 36. The method of Claim 35, wherein the cell population comprising the cultured human pluripotent cell contains at least 80% of a neural cell.
- 37. The method of Claim 32, wherein the concentration of the ceramide compound is from approximately 0.1 μ M to approximately 1000 μ M.
- 38. The method of Claim 32, wherein the concentration of the ceramide compound is from approximately 1 μ M to approximately 100 μ M.
- 39. The method of Claim 32, wherein the concentration of the ceramide compound is from approximately 5 μ M to approximately 50 μ M.
- 40. The method of Claim 32, wherein the concentration of the ceramide compound is approximately $10 \mu M$.
- 41. The method of Claim 32, wherein R= a saturated or mono- or polyunsaturated (cis or trans) alkyl group having from 12-20 carbon atoms.
- 42. The method of Claim 41, wherein the hydroxylated alkyl groups have from 1-6 carbon atoms.
- 43. The method of Claim 42, wherein R1 and R2 are hydroxylated alkyl groups.
- 44. A composition for promoting maintenance, proliferation, or differentiation of a human neural cell, the composition comprising a cell culture medium comprising MEDII conditioned medium and a ceramide compound of the general formula

wherein

R=a saturated or mono- or polyunsaturated (cis or trans) alkyl group having greater than 2 carbon atoms, and

R1, R2, R3 and R4 may be the same or different and are saturated or mono-or polyunsaturated hydroxylated alkyl groups, aryl groups, or hydrogen.

45. The composition of Claim 44, wherein the composition comprises a ceramide compound of the formula

46. The composition of Claim 44, wherein the composition comprises a ceramide compound of the formula

- 47. The composition of Claim 44, wherein the concentration of the ceramide compound is from approximately 1 μ M to approximately 100 μ M.
- 48. The composition of Claim 44, wherein the concentration of the ceramide compound is approximately 10 μ M.
- 49. The composition of Claim 44, wherein the human neural cell is cultured with the composition for a period of from approximately 6 hours to 10 days.

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50. A composition for promoting maintenance, proliferation, or differentiation of a human neural cell, the composition comprising a cell culture medium comprising MEDII conditioned medium and a sphingosine compound of the general formula

wherein

R=a saturated or mono- or polyunsaturated (cis or trans) alkyl group having greater than 2 carbon atoms, and

R1, R2, R3 and R4 may be the same or different and are saturated or mono-or polyunsaturated hydroxylated alkyl groups, aryl groups, or hydrogen.

- 51. The composition of Claim 50, wherein the concentration of the sphingosine compound is from approximately 1 μ M to approximately 100 μ M.
- 52. The composition of Claim 50, wherein the concentration of the sphingosine compound is approximately 10 μ M.
- 53. The composition of Claim 50, wherein the human neural cell is cultured with the composition for a period of from approximately 6 hours to 10 days.
- 54. A composition for promoting maintenance, proliferation, or differentiation of a human neural cell, the composition comprising a cell culture medium comprising MEDII conditioned medium and a hydroxyalkyl ester compound of the general formula

wherein

R=a saturated or mono- or polyunsaturated (cis or trans) alkyl group having greater than 2 carbon atoms, and

R1 is a saturated or mono-or polyunsaturated hydroxylated alkyl group, aryl group, or hydrogen.

55. The composition of Claim 54, wherein the composition comprises a hydroxyalkyl ester compound of the formula

- 56. The composition of Claim 55, wherein the concentration of the hydroxyalkyl ester compound is from approximately 1 μ M to approximately 100 μ M.
- 57. The composition of Claim 55, wherein the concentration of the hydroxyalkyl ester compound is approximately $10 \mu M$.
- 58. The composition of Claim 55, wherein the human neural cell is cultured with the composition for a period of from approximately 6 hours to 10 days.
- 59. A neural cell cultured in the composition of any of Claims 44-58.
- 60. A cell culture composition comprising a differentiating pluripotent human cell and a composition comprising a ceramide compound of the general formula

wherein

R=a saturated or mono- or polyunsaturated (cis or trans) alkyl group having greater than 2 carbon atoms, and

R1, R2, R3 and R4 may be the same or different and are saturated or mono-or polyunsaturated hydroxylated alkyl groups, aryl groups, or hydrogen.

61. The composition of Claim 60, wherein the composition comprises a ceramide compound of the formula

62. The composition of Claim 60, wherein the composition comprises a ceramide compound of the formula

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- 63. The composition of Claim 60, wherein the concentration of the ceramide compound is from approximately 1 μ M to approximately 100 μ M.
- 64. The composition of Claim 60, wherein the concentration of the ceramide compound is approximately $10 \mu M$.
- 65. The composition of Claim 60, wherein the differentiating pluripotent human cell is cultured with the composition for a period of from approximately 6 hours to 10 days.
- 66. The method of Claim 60, wherein R= a saturated or mono- or polyunsaturated (cis or trans) alkyl group having from 12-20 carbon atoms.
- 67. The method of Claim 66, wherein the hydroxylated alkyl groups have from 1-6 carbon atoms.
- 68. The method of Claim 67, wherein R1 and R2 are hydroxylated alkyl groups.
- 69. The method of Claim 60, wherein the pluripotent human cell is selected from the group consisting of a human embryonic stem cell, a human inner cell mass (ICM)/epiblast cell, a human primitive ectoderm cell, and a human primordial germ cell.
- 70. The method of Claim 60, wherein the pluripotent human cell is a human embryonic stem cell.
- 71. The method of Claim 60, wherein the differentiating human pluripotent cell is a neural precursor cell.